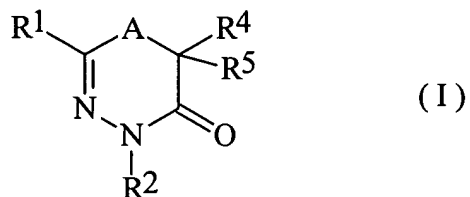


1. (Amended) A compound represented by the following formula (I), a pharmacologically acceptable salt thereof or hydrates thereof:

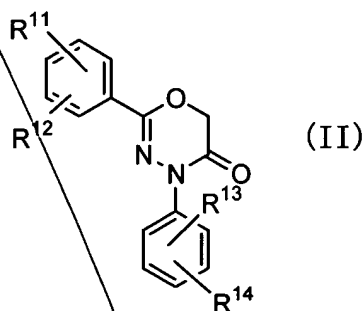


wherein A represents oxygen, sulfur or a group represented by the formula $>NR^3$ (wherein R^3 represents hydrogen atom or a lower alkyl group); R^1 represents an optionally substituted aryl group, an optionally substituted heteroaryl group that is formed from one or two 5-6 membered rings that may contain from 1 to 4 heteroatoms, an optionally substituted aralkyl group, an optionally substituted heteroaryl alkyl group, an optionally substituted aryl alkenyl group, an optionally substituted heteroaryl alkenyl group, an optionally substituted piperidyl group, an optionally substituted piperazinyl group, a morpholinyl group, an optionally substituted lower C_{3-8} cycloalkyl group, a tetrahydrofuranyl group, a tetrahydropyranyl group, an adamantyl group, an optionally substituted amino group or an optionally substituted amide group that is $-CO-N(R_a)R_b$, wherein R_a and R_b are hydrogen and C_{1-6} alkyl groups; R^2 represents an optionally substituted aryl group, an optionally substituted heteroaryl group that is formed from one or two 5-6

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membered rings that may contain from 1 to 4 heteroatoms, an optionally substituted aryl alkenyl group, an optionally substituted heteroaryl alkenyl group, an optionally substituted piperidyl group, an optionally substituted piperazinyl group, a morpholinyl group, an optionally substituted lower C₃₋₈ cycloalkyl group, a tetrahydrofuranyl group, a tetrahydropyranyl group, an adamantyl group, an optionally substituted amino group or an optionally substituted amide group that is -CO-N(R_a)R_b, wherein R_a and R_b are hydrogen and C₁₋₆ alkyl group; and R⁴ and R⁵ are the same as or different from each other and each represents hydrogen atom, hydroxyl group, nitrile group, nitro group, a lower alkyl group, an aryl group or a heteroaryl group that is formed from one or two 5 or 6 membered rings that may contain from 1 to 4 heteroatoms, provided that A is an oxygen atom, when R¹ and R² are both phenyl; and when A is a sulfur atom, R¹ is an aryl which may have a substituent, a heteroaryl which may have a substituent that is formed from one or two 5-6 membered rings that may contain 1-4 heteroatoms, an alkyl which may have a substituent, a heteroarylalkyl which may have a substituent, an arylalkenyl which may have a substituent, a heteroarylalkenyl which may have a substituent,

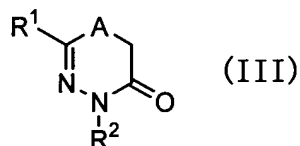
a piperidyl which may have a substituent,
 a piperadiny1 which may have a substituent,
 a morpholinyl which may have a substituent,
 a lower C₃₋₈ cycloalkyl which may have a substituent,
 tetrahydrofuranyl,
 adamantyl or
 an optionally substituted amide, that is -CO-N(R_a)R_b, wherein R_a
 and R_b are hydrogen and C₁₋₆ alkyl group; and
 provided that the compounds represented by the following
 formula (II):



(wherein R¹¹ and R¹² are the same as or different from each other
 and each represents hydrogen atom, fluorine, chlorine, bromine,
 iodine, a C1-C2 fluoroalkyl group, a C1-C2 chloroalkyl group, a
 C1-C2 bromoalkyl group, a C1-C6 alkyl group, a C3-C6 cycloalkyl
 group, a C7-C9 aralkyl group, phenyl group, a C1-C6 alkoxy
 group, a C1-C6 alkylthio group, a C1-C6 alkylsulfinyl group, a
 C7-C9 aralkoxy group, phenoxy group, phenylthio group,
 phenylsulfonyl group, an alkali metal carboxylate C2-C5
 alkoxycarbonyl group or a group represented by the formula -

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 N(R¹⁵)R¹⁶ (wherein R¹⁵ and R¹⁶ are the same as or different from each other and each represents hydrogen atom or a C1-C2 alkyl group); and R¹³ and R¹⁴ are the same as or different from each other and each represents a C₁₋₄ alkylsulfonyl group, nitro group, a group represented by the formula -OCH_nX_{3-n} (wherein X represents fluorine, chlorine, bromine or iodine; and n is an integer of 1 to 3) or the same groups as defined above for R¹¹ and R¹²) are excluded.~~

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 7. (Amended) The compound according to claim 1, wherein R⁴ and R⁵ are hydrogen and which is represented by the following formula (III):~~



~~(wherein A, R¹ and R² have the same meanings as defined in claim 1), a pharmacologically acceptable salt thereof or hydrates thereof.~~

~~8. (Amended) The compound according to claim 7, a pharmacologically acceptable salt thereof or hydrates thereof, wherein R¹ is an optionally substituted aryl group, an optionally substituted heteroaryl group that is formed from one or two 5 or 6 membered rings that may contain from 1 to 4 heteroatoms, an~~

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optionally substituted aralkyl group, an optionally substituted heteroaryl alkyl group, an optionally substituted aryl alkenyl group, an optionally substituted heteroaryl alkenyl group, a morpholinyl group, a lower C₃₋₈ cycloalkyl group, an optionally substituted amino group or an optically substituted amide group that is CO-N(R_a)R_b, wherein R_a and R_b are hydrogen and C₁₋₆ alkyl group; and R² is an optionally substituted aryl group, an optionally substituted heteroaryl group that is formed from one or two 5 or 6 membered rings that may contain from 1 to 4 heteroatoms, an optionally substituted aralkyl group, an optionally substituted heteroaryl alkyl group, a lower C₃₋₈ cycloalkyl group, a tetrahydrofuranyl group, a tetrahydropyranyl group, an optionally substituted piperidyl group or an adamantyl group.

9. (Amended) The compound according to claim 7 or 8, a pharmacologically acceptable salt thereof or hydrates thereof, wherein the substituent groups on R¹ and R² are hydrogen atom, halogen atom, hydroxyl group, lower alkyl group, lower alkenyl group, lower alkynyl group, lower alkoxy group, lower thioalkoxy group, hydroxy lower thioalkoxy group, arylthio group, heteroaryl thio group, heteroaryl(hydroxy)alkyl group, halogenated lower alkyl group, hydroxy lower alkyl group, dihydroxy lower alkyl group, halogenated (hydroxy) lower alkyl

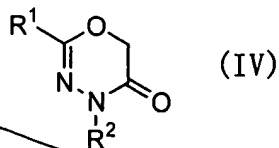
group, hydroxyalkenyl group, hydroxyalkynyl group, hydroxy lower cycloalkenyl group, lower alkoxy(hydroxy)alkyl group, lower alkoxy(hydroxy)alkoxy group, lower alkoxy alkyl group, lower alkoxy alkoxy group, lower thioalkoxy alkoxy group, lower alkyl sulfonyl alkoxy group, hydroxy lower alkoxy group, dihydroxy lower alkoxy group, hydroxy lower alkyl alkoxy group, hydroxy imino lower alkyl group, lower cycloalkyl(hydroxy)alkyl group, aralkyl group, hydroxyaralkyl group, cyano group, cyano lower alkyl group, amide group that is $-CO-N(R_a)R_b$, wherein R_a and R_b are hydrogen or C_{1-6} alkyl group, N-lower alkyl amide group, N-lower cycloalkyl amide group, N,N-di-lower alkyl amide group, N-hydroxy lower alkyl amide group, N-hydroxy lower alkyl-N-lower alkyl amide group, N-aryl amide group, cyclic aminocarbonyl group, carbamoyl group, N-lower alkyl carbamoyl group, N,N-di-lower alkyl carbamoyl group, aminosulfonyl group, cyclic aminosulfonyl group, N-lower alkyl aminosulfonyl group, N-lower cycloalkyl aminosulfonyl group, N,N-di-lower alkyl aminosulfonyl group, N-hydroxy lower alkyl aminosulfonyl group, N-lower alkoxy alkyl aminosulfonyl group, N-halogenated lower alkyl sulfonyl group, pyrrolidinyl sulfonyl group, lower alkyl sulfonyl amino alkyl group, N-lower alkyl aminosulfonyl alkyl group, N,N-di-lower alkyl aminosulfonyl alkyl group, lower acyl group, lower acyl alkyl group, lower cycloalkyl(hydroxy)methyl group, tetrahydropyranyl group, hydroxytetrahydropyranyl group, hydroxy

lower alkyl tetrahydropyranyl group, lower acyl amino alkyl group, (thiazole-2-yl)hydroxymethyl group, di(thiazole-2-yl)hydroxymethyl group, lower alkyl sulfonyl group, lower alkoxy alkyl sulfonyl group, hydroxy lower alkyl sulfonyl group, lower alkyl sulfonyl alkyl group, N-lower alkyl amide alkyl group, aryl group, aralkyl group, heteroaryl group that is formed from one or two 5 or 6 membered rings that may contain from 1 to 4 heteroatoms, heteroaryl lower alkyl group, heteroaryl lower alkoxy group, heteroaryl sulfonyl group, 4-morpholinyl sulfonyl group, 4-oxythiomorpholinyl sulfonyl group, 4-dioxythiomorpholinyl sulfonyl group, 4-morpholinyl sulfonyl group, hydroxy lower cycloalkyl group, hydroxy lower cycloalkyloxy group, hydroxy cycloalkenyl group, halogenated hydroxy lower alkyl group, 4-hydroxypiperidyl group, 4-lower alkoxypiperidyl group, ω,ω -lower alkylene dioxyalkyl group, ω,ω -lower alkylene dioxy alkoxy group, lower cycloalkyl hydroxy methyl group, aryloxy group, aryl aminosulfonyl group, amino group, lower alkyl amino group, di-lower alkyl amino group, hydroxy lower alkyl amino group, lower acyl amino group, hydroxy lower acyl amino group, lower alkyl sulfonyl amino group, pyridyl lower alkoxy group, lower alkyl pyridyl alkoxy group, lower alkoxy hydroxy alkoxy group, lower thioalkoxy alkoxy group, lower alkyl sulfonyl alkoxy group, N-lower alkyl carbamoyl group, N,N-di-lower alkyl carbamoyl group, N-hydroxy

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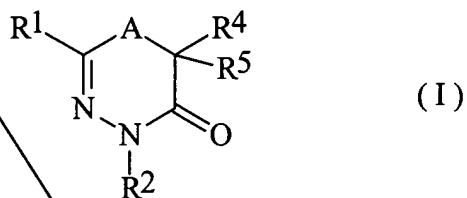
lower alkyl carbamoyl group, N-hydroxy lower alkyl-N-lower alkyl carbamoyl group, halogenated lower alkoxy group, cyano lower alkoxy group, hydroxy lower cycloalkoxy group, trifluoromethyl group, trifluoromethoxy group, amino lower alkoxy group, N-lower alkyl aminoalkoxy group, N,N-di-lower alkyl aminoalkoxy group, lower acyl alkoxy group, lower acyl aminoalkoxy group, (1,3-dioxolanyl) lower alkyl group, (1,3-dioxolanyl) lower alkoxy group, amide lower alkoxy group, 4-(hydroxy alkyl)tetrahydropyran-4-yl group, 2,3-dihydrobenzofuranyl group, 2-hydroxy-2-alkyl-2,3-dihydrobenzofuranyl group, indanonyl group, hydroxyindanyl group, imidazolyl lower alkoxy group, succimide group or 2-oxazolidone-3-yl group, optionally substituted benzoyloxy lower alkyl group, optionally substituted amino lower alkyl group, optionally substituted amino lower alkoxy group, optionally substituted aralkyloxy group, optionally substituted heteroaryl alkoxy group, optionally substituted morpholinyl lower alkoxy group, optionally substituted piperidyl lower alkoxy group, optionally substituted piperazinyl lower alkoxy group or optionally substituted pyrrolidinyl lower alkoxy group.

10. (Amended) The compound according to claim 7, represented by the following formula (IV):



(wherein R¹ and R² have the same meanings as defined in claim 7),
a pharmacologically acceptable salt thereof or hydrates thereof.

13. (Amended) A pharmaceutical composition comprising a
pharmacologically acceptable amount of the compound represented
by the following formula (I), a pharmaceutically acceptable salt
thereof or hydrates thereof, and pharmacologically acceptable
carriers:



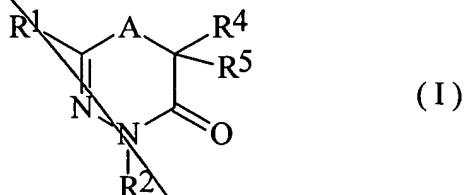
wherein A represents oxygen, sulfur or a group represented
by the formula >NR³ (wherein R³ represents hydrogen atom or a
lower alkyl group); R¹ and R² are the same as or different from
each other and each represents an optionally substituted aryl
group, an optionally substituted heteroaryl group that is formed
from one or two 5 or 6 membered rings that may contain from 1 to
4 heteroatoms, an optionally substituted aralkyl group, an
optionally substituted heteroaryl alkyl group, an optionally
substituted aryl alkenyl group, an optionally substituted

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heteroaryl alkenyl group, an optionally substituted piperidyl group, an optionally substituted piperazinyl group, a morpholinyl group, an optionally substituted lower C₃₋₈ cycloalkyl group, a tetrahydrofuranyl group, a tetrahydropyranyl group, an adamantyl group, an optionally substituted amino group or an optionally substituted amide group that is CO-N(R_a)R_b, wherein R_a and R_b are hydrogen and C₁₋₆ alkyl group; and R⁴ and R⁵ are the same as or different from each other and each represents hydrogen atom, hydroxyl group, halogen atom, nitrile group, nitro group, a lower alkyl group, an aryl group or a heteroaryl group that is formed from one or two 5 or 6 membered rings that may contain from 1 to 4 heteroatoms provided that A is an oxygen atom, when R¹ and R² are both phenyl; and when A is a sulfur atom, R¹ is an aryl which may have a substituent, a heteroaryl which may have a substituent that is formed from one or two 5-6 membered rings that may contain 1-4 heteroatoms, an aralkyl which may have a substituent, a heteroarylalkyl which may have a substituent an arylalkenyl which may have a substituent, a heteroarylalkenyl which may have a substituent, a piperidyl which may have a substituent, a piperadinyll which may have a substituent,

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a morpholinyl which may have a substituent,
a lower C₃₋₈ cycloalkyl which may have a substituent,
tetrahydrofuranyl,
adamantyl or
an optionally substituted amide, that is -CO-N(R_a)R_b, wherein R_a
and R_b are hydrogen and C₁₋₆ alkyl group.

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15. (Amended) A pharmaceutical preparation comprising the
compound represented by the following formula (I), a
pharmaceutically acceptable salt thereof or hydrates thereof:



wherein A represents oxygen, sulfur or a group represented
by the formula >NR³ (wherein R³ represents hydrogen atom or a
lower alkyl group); R¹ and R² are the same as or different from
each other and each represents an optionally substituted aryl
group, an optionally substituted heteroaryl group that is formed
from one or two 5-6 membered rings that may contain from 1 to 4
heteroatoms, an optionally substituted aralkyl group, an
optionally substituted heteroaryl alkyl group, an optionally
substituted aryl alkenyl group, an optionally substituted

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heteroaryl alkenyl group, an optionally substituted piperidyl group, an optionally substituted piperazinyl group, a morpholinyl group, an optionally substituted lower C₃₋₈ cycloalkyl group, a tetrahydrofuranyl group, a tetrahydropyranyl group, an adamantyl group, an optionally substituted amino group or an optionally substituted amide group that is -CO-N(R_a)R_b, wherein R_a and R_b are hydrogen or C₁₋₆ alkyl group; and R⁴ and R⁵ are the same as or different from each other and each represents hydrogen atom, hydroxyl group, a halogen atom, nitrile group, nitro group, a lower alkyl group, an aryl group or a heteroaryl group that is formed from one or two 5 or 6 membered rings that may contain from 1 to 4 heteroatoms provided that A is an oxygen atom, when R¹ and R² are both phenyl; and when A is a sulfur atom, R¹ is an aryl which may have a substituent, a heteroaryl which may have a substituent that is formed from one or two 5-6 membered rings that may contain 1-4 heteroatoms, an aralkyl which may have a substituent, a heteroarylalkyl which may have a substituent, an arylalkenyl which may have a substituent, a heteroarylalkenyl which may have a substituent, a piperidyl which may have a substituent, a piperadinyll which may have a substituent,

~~a morpholinyl which may have a substituent,
 a lower C₃₋₈ cycloalkyl which may have a substituent,
 tetrahydrofuranyl,
 adamantyl or
 an optionally substituted amide, that is -CO-N(Ra)Rb, wherein Ra
 and Rb are hydrogen and C₁₋₆ alkyl group.~~

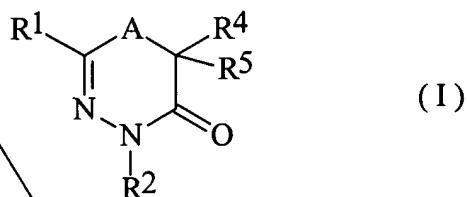
~~24. (Amended) A method of treating and ameliorating nerve
 degeneration diseases, which comprises administering a
 pharmacologically effective amount of the pharmaceutical
 preparation according to claim 15 or 16 to a patient.~~

~~25. (Amended) A method of treating and ameliorating
 demyelinating nerve diseases, which comprises administering a
 pharmacologically effective amount of the pharmaceutical
 preparation according to claim 15 or 16 to a patient.~~

~~26. (Amended) A method of treating and ameliorating acute
 nerve degeneration after cerebral ischemia, traumas in the head
 and spinal injuries, Alzheimer's disease, Parkinson's disease,
 amyotrophic lateral sclerosis, Huntington's chorea, epilepsy,
 pain, multiple sclerosis, encephalomyelitis, Guillain Barre
 syndrome, Marchiafava Bignami disease, Devic disease, Balo
 disease, HIV or HTLV myelopathy or leukoencephalopathy, which~~

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comprises administering a pharmacologically effective amount of the pharmaceutical preparation according to claim 15 or 16 to a patient.

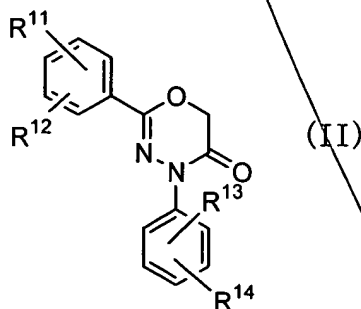
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Claim 32. A compound represented by the following formula (I), a pharmacologically acceptable salt thereof or hydrates thereof:



wherein A represents oxygen, sulfur or a group represented by the formula $>NR^3$ (wherein R^3 represents hydrogen atom or a lower alkyl group); R^1 represents an optionally substituted aryl group, an optionally substituted heteroaryl group that is formed from one or two 5 or 6 membered rings that may contain from 1 to 4 heteroatoms, an optionally substituted aralkyl group, an optionally substituted heteroaryl alkyl group, an optionally substituted aryl alkenyl group, an optionally substituted heteroaryl alkenyl group, an optionally substituted piperidyl group, an optionally substituted piperazinyl group, a morpholinyl group, an optionally substituted lower C_{3-8} cycloalkyl group, a tetrahydrofuranyl group, a tetrahydropyranyl

group, an adamantyl group, an optionally substituted amino group or an optionally substituted amide group that is $-\text{CO}-\text{N}(\text{R}_a)\text{R}_b$ wherein R_a and R_b are hydrogen and C_{1-6} alkyl group; R^2 represents an optionally substituted aryl group, an optionally substituted heteroaryl group that is formed from one or two 5 or 6 membered rings that may contain from 1 to 4 heteroatoms, an optionally substituted aralkyl group wherein it is not benzyl, an optionally substituted heteroarylalkyl group wherein it is not pyrimidinyl alkyl, an optionally substituted aryl alkenyl group, an optionally substituted heteroaryl alkenyl group, an optionally substituted piperidyl group, an optionally substituted piperazinyl group, a morpholinyl group, an optionally substituted lower C_{3-8} cycloalkyl group, a tetrahydrofuranyl group, a tetrahydropyranyl group, an adamantyl group, an optionally substituted amino group or an optionally substituted amide group that is $-\text{CO}-\text{N}(\text{R}_a)\text{R}_b$, wherein R_a and R_b are hydrogen and C_{1-6} alkyl group; and R^4 and R^5 are the same as or different from each other and each represents hydrogen atom, hydroxyl group, nitrile group, nitro group, a lower alkyl group, an aryl group or a heteroaryl group that is formed from one or two 5 or 6 membered rings that may contain from 1 to 4 heteroatoms, provided that A is an oxygen atom, when R^1 and R^2 are both phenyl; and

when A is a sulfur atom, R¹ is
 an aryl which may have a substituent,
 a heteroaryl which may have a substituent that is formed from
 one or two 5-6 membered rings that may contain 1-4 heteroatoms,
 an aralkyl which may have a substituent,
 a heteroarylalkyl which may have a substituent,
 an arylalkenyl which may have a substituent,
 a heteroarylalkenyl which may have a substituent,
 a piperidyl which may have a substituent,
 a piperadinyll which may have a substituent,
 a morpholinyl which may have a substituent,
 a lower C₃₋₈ cycloalkyl which may have a substituent,
 tetrahydrofuranyl,
 adamantyl or
 an optionally substituted amide, that is -CO-N(R_a)R_b, wherein R_a
 and R_b are hydrogen and C₁₋₆ alkyl group; and
 provided that the compounds represented by the following
 formula (II):



~~(wherein, R^{11} and R^{12} are the same as or different from each other and each represents hydrogen atom, fluorine, chlorine, bromine, iodine, a C1-C2 fluoroalkyl group, a C1-C2 chloroalkyl group, a C1-C2 bromoalkyl group, a C1-C6 alkyl group, a C3-C6 cycloalkyl group, a C7-C9 aralkyl group, phenyl group, a C1-C6 alkoxy group, a C1-C6 alkylthio group, a C1-C6 alkylsulfinyl group, a C7-C9 aralkoxy group, phenoxy group, phenylthio group, phenylsulfonyl group, an alkali metal carboxylate C2-C5 alkoxy carbonyl group or a group represented by the formula $-N(R^{15})R^{16}$ (wherein R^{15} and R^{16} are the same as or different from each other and each represents hydrogen atom or a C1-C2 alkyl group); and R^{13} and R^{14} are the same as or different from each other and each represents a C₁₋₄ alkylsulfonyl group, nitro group, a group represented by the formula $-OCH_nX_{3-n}$ (wherein X represents fluorine, chlorine, bromine or iodine; and n is an integer of 1 to 3) or the same groups as defined above for R^{11} and R^{12}) are excluded.~~

Please add the following claims:

~~--33. (new) A method of preventing a disease selected from the group consisting of epilepsy, multiple sclerosis, Huntington's chorea, Alzheimer's disease, Parkinson's disease and amyotrophic lateral sclerosis comprising administering an effective amount of~~